

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-23-1243; Docket No. CDC-2022-0134]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC),
Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

summary: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other federal agencies the opportunity to comment on a continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Rapid Response Suicide Investigation Data Collection. This data collection is designed to inform the implementation of prevention strategies in a state, county, community, or vulnerable population where a possible suicide cluster or increasing trend has been observed.

DATES: CDC must receive written comments on or before [INSERT DATE 60 DAYS AFTER PUBLICATION DATE IN THE FEDERAL REGISTER].

ADDRESSES: You may submit comments, identified by Docket No.

CDC-2022-0134 by either of the following methods:

- Federal eRulemaking Portal: www.regulations.gov. Follow the instructions for submitting comments.
- Mail: Jeffrey M. Zirger, Information Collection Review Office,
 Centers for Disease Control and Prevention, 1600 Clifton Road,
 NE, MS H21-8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency
name and Docket Number. CDC will post, without change, all
relevant comments to www.regulations.gov.

Please note: Submit all comments through the Federal eRulemaking

portal (www.regulations.gov) or by U.S. mail to the address

listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road, NE, MS H21-8, Atlanta, Georgia 30329; Telephone: 404-639-7570; E-mail: omb@cdc.gov.

SUPPLEMENTARY INFORMATION:

Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires federal agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each new proposed collection, each

proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

- 1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- 2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- 3. Enhance the quality, utility, and clarity of the information to be collected;
- 4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses; and
- 5. Assess information collection costs.

Proposed Project

Rapid Response Suicide Investigation Data Collection (OMB Control No. 0920-1243, Exp. 5/31/2021) - Extension - National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC is frequently called upon to respond to urgent requests from one or more external partners (e.g., local, state, territory, and tribal health authorities; other federal agencies; local and state leaders; schools; or other partner organizations) to conduct investigations of suicide. Supporting rapid investigations to inform the implementation of effective suicide prevention strategies is one of the most important ways CDC can serve to protect and promote the health of the public.

Rapid Response Suicide Investigation Data Collections are specifically designed to inform the implementation of prevention strategies in a state, county, community, or vulnerable population where a possible suicide cluster or increasing trend has been observed. This generic clearance will not be used to conduct research studies or to collect data designed to draw conclusions about the United States or areas beyond the defined geographic location or vulnerable population that is the focus of the investigation. CDC in collaboration with external partners (e.g., local, state, territory, and tribal health authorities; other federal agencies; local and state leaders; schools; or other partner organizations) will identify the

respondent universe for each Rapid Response Suicide

Investigation Data Collection. The respondent universe will be determined based on the information needed to understand potential suicide clusters, significant increases in suicidal behavior and suicide, risk and protective factors, and vulnerable populations in order to inform the implementation of suicide prevention strategies. When the goal is generalizability, CDC will submit the sampling methods to OMB as part of the GenIC package.

CDC requests OMB approval for an estimated 1,000 annual burden hours. There are no costs to respondents other than their time to participate.

Estimated Annualized Burden Hours

Type of Respondent	Form Name	Number of Respondents	Number of Responses per respondent	Average Burden per Response (in hours)	Total Burden Hours
Rapid Response Suicide Investigation Data Collection Participants	Rapid Response Suicide Investigation Protocol	2,000	1	30/60	1,000

Jeffrey M. Zirger,

Lead,

Information Collection Review Office,

Office of Scientific Integrity,
Office of Science,
Centers for Disease Control and Prevention.

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